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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO
09/813,444	03/20/2001	Brent Iverson	MXGN:005USC2	3269
7590 01/04/2006			EXAMINER	
Steven L. Highlander, Esq. FULBRIGHT & JAWORSKI L.L.P.			DO, PENSEE T	
Suite 2400	E JAWORSKI L.L.P.	ART UNIT	PAPER NUMBER	
600 Congress Avenue Austin, TX 78701			1641	
			DATE MAILED: 01/04/2000	5

Please find below and/or attached an Office communication concerning this application or proceeding.

		Applic	ation No.	Applicant(s)				
Cifice Action Summary		09/81	3,444	IVERSON ET AL.				
		Exami	ner	Art Unit				
		Pense	e T. Do	1641				
Period fo	The MAILING DATE of this commun or Reply	ication appears on	the cover sheet	with the correspondence ad	Idress			
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Status					,			
1)🛛	Responsive to communication(s) file	d on <i>14 July 200</i> 5	i .					
	This action is FINAL . 2b) This action is non-final.							
3)□	Since this application is in condition for allowance except for formal matters, prosecution as to the merits is							
,—	closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213.							
Disposit	ion of Claims							
4)⊠	4)⊠ Claim(s) <u>1-3,6-12,15-26 and 46</u> is/are pending in the application.							
	4a) Of the above claim(s) is/are withdrawn from consideration.							
5)□)☐ Claim(s) is/are allowed.							
6)⊠	Claim(s) <u>1-3,6,11,15, 16 and 46</u> is/are rejected.							
7)	Claim(s) <u>7-10,12 and 17-26</u> is/are objected to.							
8)□	Claim(s) are subject to restrict	tion and/or electio	n requirement.					
Applicati	ion Papers							
9)	The specification is objected to by the	e Examiner.						
10)	10)☐ The drawing(s) filed on is/are: a)☐ accepted or b)☐ objected to by the Examiner.							
	Applicant may not request that any object	ction to the drawing(s) be held in abey	ance. See 37 CFR 1.85(a).				
	Replacement drawing sheet(s) including		•		` '			
11)	The oath or declaration is objected to	by the Examiner.	Note the attach	ed Office Action or form P1	Г О-152 .			
Priority (ınder 35 U.S.C. § 119							
12)	Acknowledgment is made of a claim	for foreign priority	under 35 U.S.C.	. § 119(a)-(d) or (f).				
	a) ☐ All b) ☐ Some * c) ☐ None of:							
	1. Certified copies of the priority	documents have t	een received.					
	2. Certified copies of the priority	documents have t	een received in	Application No				
	3. Copies of the certified copies	of the priority docu	iments have bee	en received in this National	Stage			
	application from the Internatio	nal Bureau (PCT I	Rule 17.2(a)).					
* 5	See the attached detailed Office action	n for a list of the c	ertified copies no	ot received.				
Attachmen	• •							
1) Notic	e of References Cited (PTO-892) e of Draftsperson's Patent Drawing Review (P	TO 048)		v Summary (PTO-413) o(s)/Mail Date				
3) 🛛 Inforr	e of Dransperson's Patent Drawing Review (P mation Disclosure Statement(s) (PTO-1449 or r No(s)/Mail Date <u>7/14/05</u> .			nformal Patent Application (PTO-152)				

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DETAILED ACTION

Amendment Entry & Claim Status

The amendment filed on July 14, 2005 has been acknowledged and entered.

Claims 1-3, 612, 15-26, and 46 are pending

Withdrawn Rejection(s)

Rejection under 112, 2nd paragraph is withdrawn herein.

Rejection under 102 (e) by Georgiou '867, Georgiou '344, and Higuchi '613 are withdrawn herein.

Maintained Rejection(s)

Claim Rejections - 35 USC § 102

The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless -

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

Claims 1-3, 6-9, 11, 15-20, 25, 26 and 46 are rejected under 35 U.S.C. 102(b) as being anticipated by Slamon et al. (US 4,918,162).

Slamon teaches methods for identifying and monitoring human cancers. The methods rely on the detection of N-myc protein in a biological specimen, usually a cell sample such as tissue sample or sputum sample. Presence of the N-myc protein in the biological specimen may be diagnostic and/or prognostic of the cancer. Polypeptides and antibodies are used for detecting the N-myc proteins, where the polypeptides are associated with immunogenic sites on the protein. The polypeptides may be natural or

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synthetic. Such polypeptides include the N-myc protein in substantially pure form as well as fragments thereof. Monoclonal or polyclonal antibodies against the polypeptides are prepared by conventional techniques. Six polypeptides capable of eliciting antibodies useful in the present method have been identified. The method of synthesizing the polypeptides involves the expression in cultured cells of recombinant DNA molecules encoding a desired portion of the N-myc gene. Suitable cDNA and genomic libraries may be obtained from human cell lines known to contain the N-myc gene. (see col. 1, line 65-col. 2, line 48; col. 4, lines 36-49). The natural or synthetic DNA fragments coding for a desired N-myc fragment will be incorporated in DNA constructs capable of introduction to and expression in an in vitro cell culture. Usually, the DNA constructs will be suitable for replication in a unicellular host, such as yeast or bacteria i.e. negative bacteria E.coli. but may also be intended for introduction and integration within the genome of cultured mammalian or other eukaryotic cell lines. DNA constructs prepared for introduction into bacteria or yeast will include a replication system recognized by the host. Available expression vectors, which include the replication system and transcriptional and translational regulatory sequences together with an insertion site for the N-myc DNA sequence may be employed. (see col. 4,lines 62-68; col. 5, lines 1-15; col. 9, lines 65-66). The polypeptide can be an antibody or antibody fragment. The step of selecting a host cell that expresses the desired polypeptides comprises the steps of contacting said antibody or antibody-fragmentexpressing cells with a selected antigen; and identifying a host cell that binds to said selected antigen (see col. 9, line 23-col. 10, line 14). The antigen/polypeptide is labeled

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with a fluorescers, chemiluminescers, magnetic particles etc. (see col. 6, lines 58-68). The vector library is obtained by administering to an animal such as a mouse a desired antigen. The mouse is then killed, the spleen removed, and the spleen cells immortalized. DNA segments that encode distinct antibodies or antibody fragments were obtained and incorporated into a plurality of expression vectors, the vectors expressing antibodies or antibody fragments on the outer membrane surface of a Gram negative host cell, E. coli. (see col. 4, lines 62-68; col. 5, line 1-68). Selected cells that express a desired antibody are subjected to cleavage to release the selected antibody or antibody fragment from the surface of the outer membrane. (see col. 7, lines 27-50). Since the reference teaches eukaryotic cell, it is inherent that insect cell is included because insect cell is a eukaryotic cell.

Response to Arguments

Applicant's arguments filed July 14, 2005 have been fully considered but they are not persuasive.

Regarding the rejection by Slamon '162, Applicants argue that Slamon fails to teach a "library" of "vectors" or candidate polypeptides. Rather the reference teaches merely the recombinant expression of a few specific N-myc proteins. Second, there is no teaching of cell surface expression. The passage to which the examiner cites teaches away from cell surface expression, requiring release of the target either by "shed" of the proteins or "lysing" of cells. (col. 7, lines 27-36).

Slamon teaches "the vector library is obtained by administering to an animal such as a mouse a desired antigen". (see col. 4, lines 62-68; col. 5, lines 1-68). Selected cells

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that express a desired antibody are subjected to cleavage to release the selected antibody or antibody fragments from the surface of the outer membrane. (see col. 7, lines 27-50). Applicants assert that the passage on col. 7, lines 27-36 teaches away from the present invention by requiring a "releasing" step of the antibody/fragment. However, such step is not excluded by the present invention because the claims of the present invention contain opening claim language which encompasses extra step such as releasing the antibody/fragments after the host cell expresses the desired antibody/fragment. Surface expression is taught on col. 4, lines 62-68).

Allowable Subject Matter

Claims 10, 12, 21-24 are objected to as being dependent upon a rejected base claim, but would be allowable if rewritten in independent form including all of the limitations of the base claim and any intervening claims.

Conclusion

Applicant's amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP § 706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any

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extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Pensee T. Do whose telephone number is 571-272-0819. The examiner can normally be reached on Monday-Friday, 7:00-3:00.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Long Le can be reached on 571-272-0823. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

Pensee T. Do Patent Examiner September 28, 2005

> LONG V. LE SUPERVISORY PATENT EXAMINER TECHNOLOGY CENTER 1600

12/27/28